

# AIRS CO2 Retrievals Using the Method of Vanishing Partial Derivatives (VPD)

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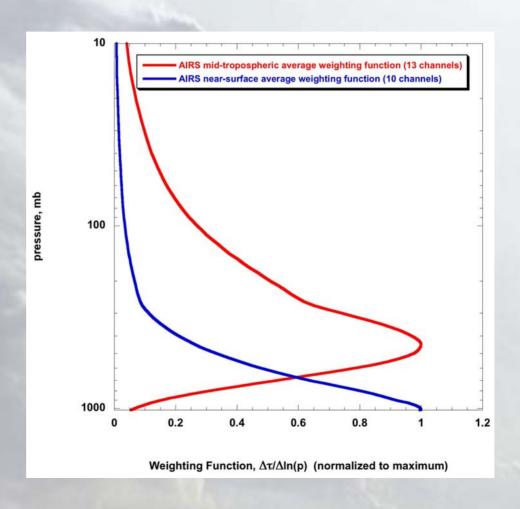
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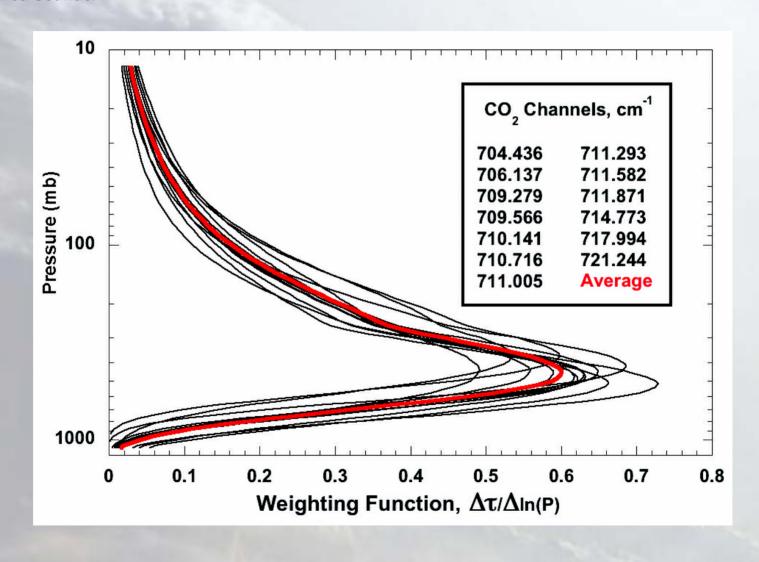


## AIRS CO<sub>2</sub> Sounding Channels Average Weighting Functions





# **CO<sub>2</sub> Sounding Channels Individual Weighting Functions**





## Paper in GRL (November 18, 2005)

doi:10.1029/2005GL024165

GEOPHYSICAL RESEARCH LETTERS, VOL. 32, L22803, doi:10.1029/2005GL024165, 2005

## On the determination of atmospheric minor gases by the method of vanishing partial derivatives with application to CO<sub>2</sub>

M. Chahine, <sup>1</sup> C. Barnet, <sup>2</sup> E. T. Olsen, <sup>1</sup> L. Chen, <sup>1</sup> and E. Maddy<sup>3</sup>

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[1] We present a general method for the determination of minor gases in the troposphere from high spectral resolution observations. In this method, we make use of a general property of the total differential of multi-variable functions to separate the contributions of each individual minor gas. We have applied this method to derive the mixing ratio of carbon dioxide in the mid-troposphere using data from the Atmospheric Infrared Sounder (AIRS) currently flying on the NASA Aqua Mission. We compare our results to the aircraft flask CO2 measurements obtained by H. Matsueda et al, over the western Pacific and demonstrate skill in tracking the measured 5 ppmv seasonal variation with an accuracy of  $0.43 \pm 1.20$  ppmv. Citation: Chahine, M., C. Barnet, E. T. Olsen, L. Chen, and E. Maddy (2005), On the determination of atmospheric minor gases by the method of vanishing partial derivatives with application to CO<sub>2</sub>, Geophys. Res. Lett., 32, L22803, doi:10.1029/ 2005GL024165.

#### 1. Introduction

#### 2. General Approach

[3] We consider the radiative transfer equation

$$R(\nu) = S_s(\nu, \varepsilon_s, \ldots) + \int_{p_s}^{0} B[\nu, T(p)] \left( \frac{\partial \tau(\nu, p, \langle \ldots \rangle)}{\partial p} \right) dp \quad (1)$$

where  $R(\nu)$ , the outgoing radiance at frequency  $\nu$  measured at the satellite, is the sum of emissions from the surface and the atmosphere. Here  $\varepsilon_s$  is the surface emissivity, B the Planck blackbody function,  $\tau$  the transmission function from any pressure level p to the top of the atmosphere and the angle bracket  $\langle \ldots \rangle$  denotes a function of the profiles of temperature T(p), water vapor q(p), ozone  $O_3(p)$ , carbon dioxide mixing ratio  $CO_2(p)$ , etc. In this paper, we express the outgoing radiance  $R(\nu)$  in brightness temperature units,  $\Theta(\nu)$ , in order to simplify its use across a wide range of frequencies.



#### **RECENT PAPERS**

Barnet, C.D., M. Goldberg, L. McMillin and M.T. Chahine (2004), Remote sounding of trace gases with the EOS/AIRS instrument, *Proceedings of SPIE*, 5548, 300-312.

Chedin, A., R. Saunders, A. Hollingsworth, N.A. Scott, M. Matricardi, J. Etcheto, C. Clerbaux, R. Armante and C. Crevoisier (2003), The feasibility of monitoring CO2 from high-resolution infrared sounders. *J. Geophys. Res.* 108, 4064-4071, doi:10.1029/2001JD001443.

Crevoisier, C., A. Chedin and N.A. Scott (2003), AIRS channel selection for CO2 and other trace-gas retrievals, *Q. J. R. Meteorol. Soc.* 129, 2719-2740.

Crevoisier, C., S. Heilliette, A. Chedin, S. Serrar, R. Armante and N.A. Scott (2004), Midtropospheric CO2 concentration retrieval from AIRS observations in the tropics, *Geophys. Res. Lett.* 31, 17106-17110, doi:10.1029/2004GL020141.

Engelen, R.J. and G.L. Stephens (2004a), Information content of infrared satellite sounding measurements with respect to CO2, *J. Appl. Meteor.* 43, 373-378.

Engelen, R.J., E. Andersson, F. Chevallier, A. Hollingsworth, M. Matricardi, A. P. McNally, J.-N. Thépaut, and P. D. Watts (2004b), Estimating atmospheric CO2 from advanced infrared satellite radiances within an operational 4D-Var data assimilation system: Methodology and first results. *J. Geophys. Res.*, 109, D19309, doi:10.1029/2004JD004777.

Engelen, R.J. and A. P. McNally (2005), Estimating atmospheric CO2 from advanced infrared satellite radiances within an operational 4D-Var data assimilation system: Results and validation. *J. Geophys. Res.* (in press), doi:10.1029/2005JD005982.

#### **Method of Vanishing Partial Derivatives (VPD)**

#### In GRL, November 18, 2005

We consider the case where observations are made in a spectral region in the infrared where several minor gases such as  $CO_2$ ,  $O_3$ , CO,  $CH_4$  and  $SO_2$  are radiatively active.

We define the residual function G as

$$G^{(n)} = \sum \left[\Theta_M(\nu) - \Theta_C^{(n)}(\nu)\right]^2$$

We aim to find the set of  $X_i^{\nu}$  which minimizes the residual function. We express the total differential of  $G_i^{\nu}$  as

$$dG = \frac{\partial G}{\partial X_1} dX_1 + \frac{\partial G}{\partial X_2} dX_2 + \frac{\partial G}{\partial X_3} dX_3 + \dots + \frac{\partial G}{\partial X_i} dX_i + \varepsilon$$
 (3)

From the general property of total differentials, the condition that G in equation (4)

should have a maximum or a minimum at a point (  $\overline{X}_i^{(1)}$  ,  $\overline{X}_2^{(1)}$  ,  $\overline{X}_i^{(1)}$  ,  $\overline{X}_i^{(1)}$  ) is that

each of the first partial derivatives should individually vanish at that point.

$$\frac{\partial G}{\partial X_1}, \frac{\partial G}{\partial X_2}, \frac{\partial G}{\partial X_3}, \dots, \frac{\partial G}{\partial X_i}$$
 (4)

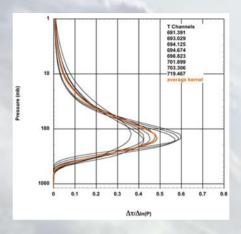
Thus we reach an important conclusion that the value of the individual mixing ratio of each of the minor gases considered is determined by the value that makes their first partial derivative in equation (4) vanish individually. Therefore, even though the observed spectra cannot differentiate between the individual lines, the partial differentials can!



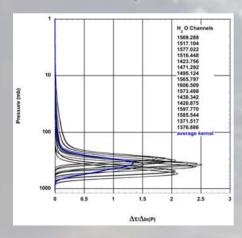
# **Auxiliary Sounding Channels Individual Weighting Functions**

#### Atmospheric Infrared Sounder

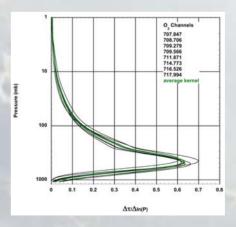
#### **Temperature**



#### Water vapor

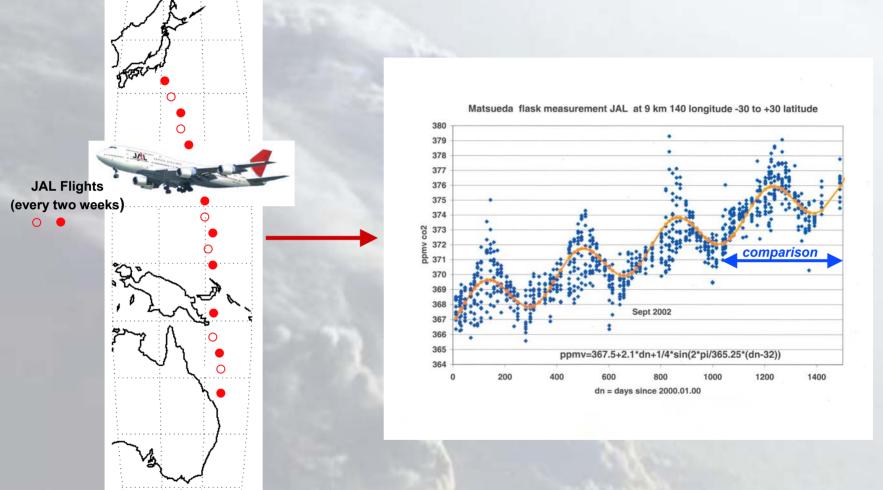


#### **Ozone**



### **Matsueda Airborne Flask**

CO<sub>2</sub> Measurements at 10.5 km altitude

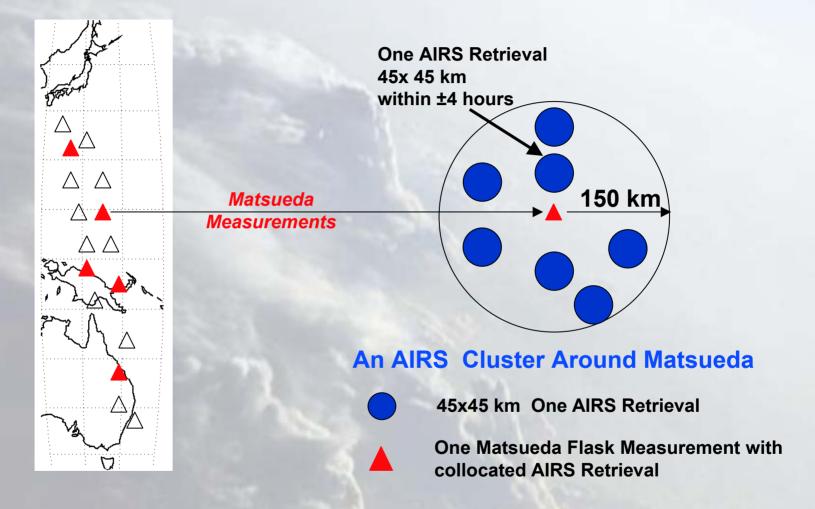


#### **Matsueda Airborne Flask**

CO<sub>2</sub> Measurements at 10.5 km altitude

#### **Collocated AIRS**

CO<sub>2</sub> Retrievals at 5-15 km altitude



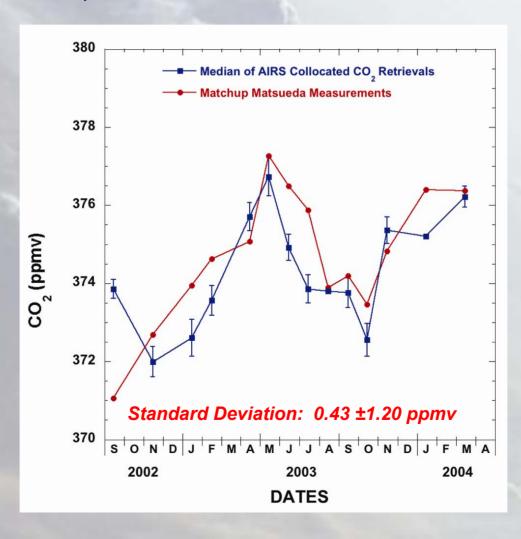
A Total of 2332 AIRS Retrievals (in the presence of clouds) found Collocated with 223 Matsueda measurements



#### Validation of AIRS CO2 Results

Comparison of AIRS CO<sub>2</sub> Retrievals with Collocated Matsueda Flask Measurements

(Collocation Criteria: ± 4 hours and 150 km radius around Matsueda)





**Starting Number of** Clusters of ~210

#### Independence of the solution

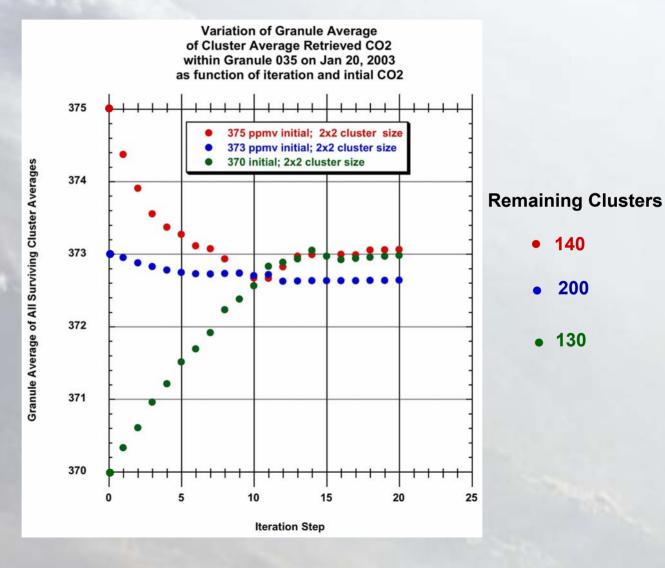
from Initial starting value of the CO2 Mixing Ratio

370 373 375 *ppmv* 

140

200

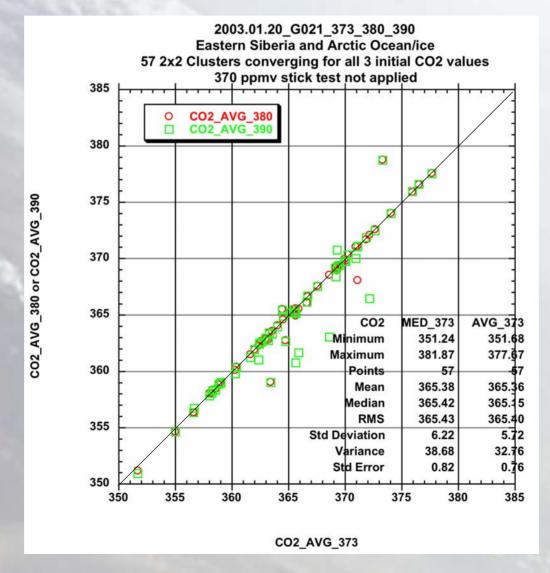
130





## Independence of the solution from Initial starting value of the CO2 Mixing Ratio

#### **Atmospheric Infrared Sounder**

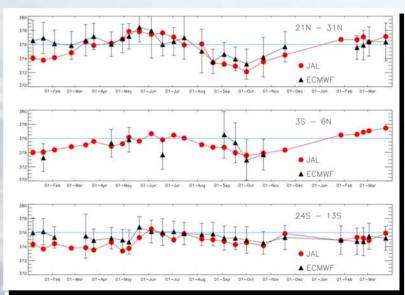




# Estimating atmospheric CO<sub>2</sub> from advanced infrared radiances with an operational 4D-Var data assimilation system: Results and Validation

#### Richard J. Engelen and Anthony P. McNally

More than a year of AIRS radiance observations have been assim-Abstract. ilated in the ECMWF 4D-Var data assimilation system to estimate tropospheric CO<sub>2</sub>. The assimilation of a set of eighteen spectral channels provides a mean tropospheric mixing ratio representing a layer between about 700 hPa and the tropopause. Analysis errors for a 5-day mean on a 6° by 6° averaging grid box are on the order of 1%. Comparisons with independent flight data from JAL and NOAA/CMDL are favourable. Differences between the averaged assimilation estimates and the onboard flask observations are generally within the 1- $\sigma$  error bars. Currently, this work is being extended by introducing CO<sub>2</sub> as a full assimilation model tracer variable, which will allow the operational monitoring of atmospheric CO<sub>2</sub> using AIRS observations and observations from upcoming instruments.



**Figure 6.** Comparison of CO<sub>2</sub> estimates with JAL observations for three different latitude zones from January 2003 to March 2004. Missing ECMWF data are caused by extensive cloud cover in the area.



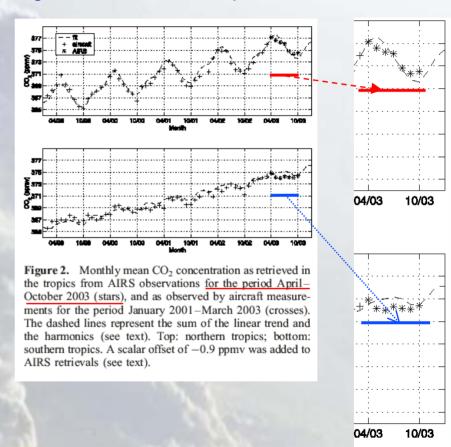
## Midtropospheric CO<sub>2</sub> concentration retrieval from AIRS observations in the tropics

Crevoisier, S Heilliette, A. Chedin, S. Serrar, R. Armante and N.A. Scott

(Using Neural networks method)

**Atmospheric Infrared Sounder** 

[1] Midtropospheric carbon dioxide (CO<sub>2</sub>) concentration is retrieved in the tropics [20S:20N], over sea, at night, for the period April to October 2003 from the Atmospheric Infrared Sounder (AIRS) observations. The method relies on a non-linear regression inference scheme using neural networks. A rough estimate of the mean precision of the method is about 2.5 ppmv (0.7%). The retrieved seasonal cycle and its latitudinal dependence agree well with aircraft CO<sub>2</sub> in situ measurements made at the same altitude range. Maps produced on a monthly basis at a resolution of 15° × 15°, although not yet fully understood, show good agreement with known characteristics of CO<sub>2</sub> distribution reflecting both atmospheric transport and surface fluxes

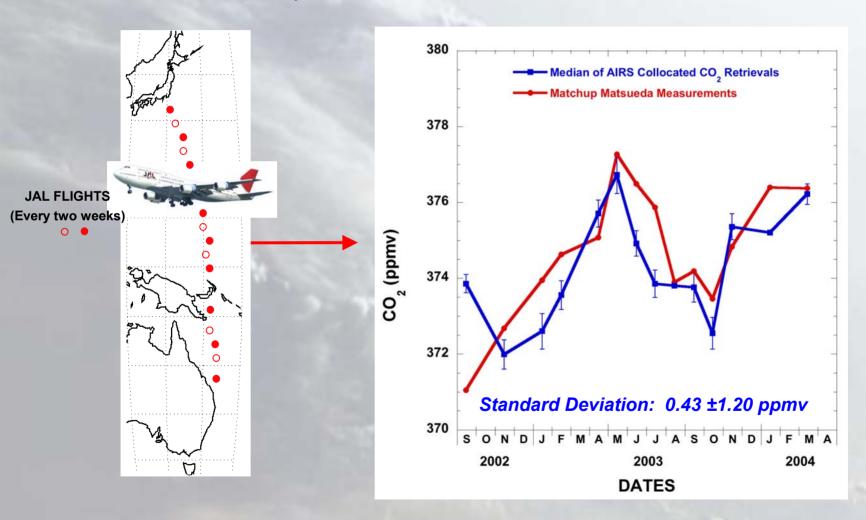




#### **SUMMARY**

Comparison of AIRS CO<sub>2</sub> Retrievals with collocated Matsueda Flask Measurements

(Colocation Criteria: ± 4 hours and 150 km radius around Matsueda)



**NEXT: GENERATE GLOBAL MAPS OF CO2 DISTRIBUTION** 



# Average CO2 AIRS Data, Jan 18-21, 2003 parts per million by volume (PPMV) 366 368 370 372 374

January 18-21 May 6-19 October 19-25

# AIRS Global CO<sub>2</sub> Maps 2003

